

Prisca 5.1.0.17
Date of report: 24/09/24

N A

Patient data			
Name Mrs. SARIKA RAMCHANDRA FARADE		Patient ID 0742409200051	
Birthday 21/12/89		Sample ID A0467505	
Age at sample date 34.7		Sample Date 20/09/24	
Gestational age 13 + 0			
Correction factors			
Fetuses 1	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 46	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 13 + 0
PAPP-A	4.36 mIU/mL	0.67	Method CRL Robinson
fb-hCG	58.2 ng/mL	1.42	Scan date 20/09/24
Risks at sampling date			Crown rump length in mm 69
Age risk 1:292			Nuchal translucency MoM 1.32
Biochemical T21 risk 1:325			Nasal bone unknown
Combined trisomy 21 risk 1:474			Sonographer N A
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT MD
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 474 women with the same data, there is one woman with a trisomy 21 pregnancy and 473 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!			
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

below cut off	Below Cut Off, but above Age Risk	above cut off
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